

## EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	450	(548/541).CCLS.	US-PGP UB; USPAT; USOCR ; EPO; JPO; DERWE NT; IBM_T DB	OR	OFF	2007/06/24 08:32
L2	160	(549/15).CCLS.	US-PGP UB; USPAT; USOCR ; EPO; JPO; DERWE NT; IBM_T DB	OR	OFF	2007/06/24 08:32
L3	0	I1 and I2	US-PGP UB; USPAT; USOCR ; EPO; JPO; DERWE NT; IBM_T DB	OR	ON	2007/06/24 08:33

10541607C>

06/24/2007

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FILE 'REGISTRY' ENTERED AT 11:10:47 ON 19 FEB 2007  
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STRUCTURE FILE UPDATES: 18 FEB 2007 HIGHEST RN 921759-52-6  
DICTIONARY FILE UPDATES: 18 FEB 2007 HIGHEST RN 921759-52-6

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TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

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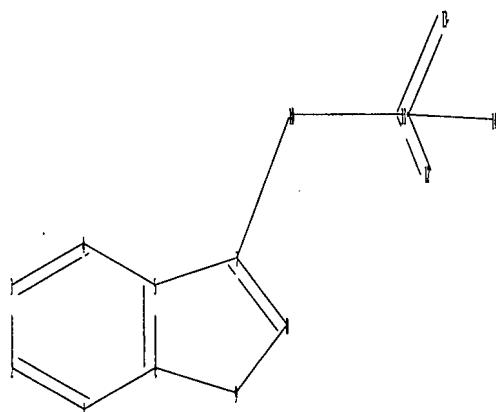
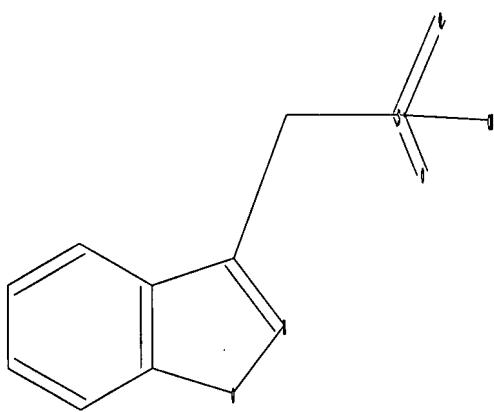
REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/reqprops.html>

=> Uploading C:\Program Files\Stnexp\Queries\10541607.str

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06/24/2007



chain nodes :  
10 11 12 13 14

ring nodes :

1 2 3 4 5 6 7 8 9

chain bonds :

7-10 10-11 11-12 11-13 11-14

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9

exact/norm bonds :

5-7 6-9 7-8 8-9 10-11

exact bonds :

7-10

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 11-12 11-13 11-14

Match level :

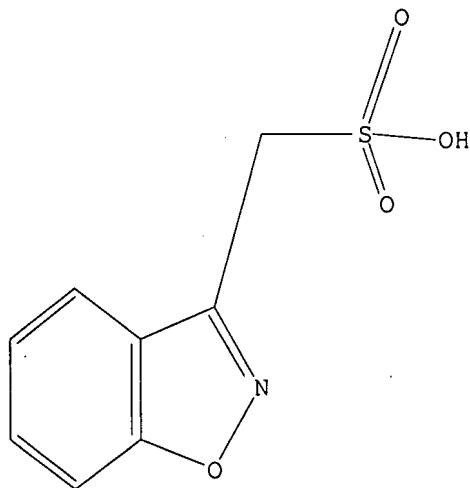
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS  
11:CLASS 12:CLASS 13:CLASS 14:CLASS

L1 STRUCTURE UPLOADED

=> D

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=>

S L1

SAMPLE SEARCH INITIATED 11:11:07 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED - 5 TO ITERATE

100.0% PROCESSED 5 ITERATIONS 0 ANSWERS  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*  
PROJECTED ITERATIONS: 5 TO 234  
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> S L1 FULL  
FULL SEARCH INITIATED 11:11:31 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 102 TO ITERATE

100.0% PROCESSED 102 ITERATIONS 14 ANSWERS  
SEARCH TIME: 00.00.01

L3 14 SEA SSS FUL L1

=> FILE CAPLUS  
COST IN U.S. DOLLARS SINCE FILE TOTAL  
SESSION  
FULL ESTIMATED COST ENTRY 172.10 172.31

FILE 'CAPLUS' ENTERED AT 11:11:36 ON 19 FEB 2007  
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FILE COVERS 1907 - 19 Feb 2007 VOL 146 ISS 9  
FILE LAST UPDATED: 18 Feb 2007 (20070218/ED)

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=> S L3  
L4 17 L3

=> S L3 AND OXIME  
17 L3  
45138 OXIME  
16097 OXIMES  
50073 OXIME  
(OXIME OR OXIMES)  
L5 2 L3 AND OXIME

=> D IBIB ABS HITSTR TOT

LS ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:606452 CAPLUS

DOCUMENT NUMBER: 141:140420

TITLE: A process for the preparation of

benzo[d]isoxazol-3-yl-

methanesulfonic acid

INVENTOR(S): Razzetti, Gabriele; Mantegazza, Simone; Castaldi, Graziano; Allegri, Pietro; Lucchini, Vittorio; Bologna, Alberto

PATENT ASSIGNEE(S): Dinamite Dipharm S.P.A., Italy

SOURCE: PCT Int. Appl., 22 pp.

CODEN: PIXKD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

WO 2004063173 A1 20040729 WO 2003-EP14919 20031224

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MM, MZ, SD, SL, SZ, T2, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG

CA 2512791 A1 20040729 CA 2003-2512791 20031224

AU 2003298248 A1 20040810 AU 2003-298248 20031224

EP 1581508 A1 20051005 EP 2003-795972 20031224

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK

JP 2006515322 T 20060525 JP 2005-512865 20031224

US 2006135582 A1 20060622 US 2005-541607 20050707

PRIORITY APPLN. INFO.: IT 2003-MI26 A 20030110

IT 2003-MI1383 A 20030704

WO 2003-EP14919 W 20031224

OTHER SOURCE(S): CASREACT 141:140420

AB The title compound (I) or its salt, useful as an intermediate in the preparation

of anticonvulsant zonisamide, is prepared by reaction of 1,2-benzoxathin-4(3H)-one 2,2-dioxide oxime (II) with organic base or alkali or alkaline earth hydroxide. Thus, reaction of II with eq NaOH at

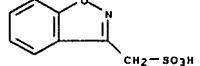
room temperature for 3 h gave 70% sodium salt of I.

IT 726188-85-8P

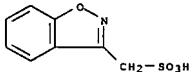
RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation) (preparation of 1,2-benzisoxazole-3-methanesulfonic acid or its salt

as

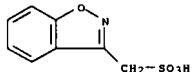
LS ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)



● Li



IT 73101-64-1P 726188-84-7P  
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
(preparation of 1,2-benzisoxazole-3-methanesulfonic acid or its salt)  
as intermediate for zonisamide;  
RN 73101-64-1 CAPLUS  
CN 1,2-Benzisoxazole-3-methanesulfonic acid, sodium salt (9CI) (CA INDEX NAME)



RN 726188-84-7 CAPLUS  
CN 1,2-Benzisoxazole-3-methanesulfonic acid, lithium salt (9CI) (CA INDEX NAME)

LS ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

NAME: 2002:695963 CAPLUS

ACCESSION NUMBER: 137:216942

TITLE: Process for the preparation of 1,2-benzisoxazole-3-acetic acid, an intermediate in the synthesis of zonisamide

INVENTOR(S): Mendelovitch, Mariyana; Nidam, Tamar; Teva Pharmaceutical Industries Ltd., Israel; Teva Pharmaceuticals USA, Inc.

PATENT ASSIGNEE(S): Teva Pharmaceutical Industries Ltd., Israel; Teva Pharmaceuticals USA, Inc.

SOURCE: PCT Int. Appl., 14 pp.

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

WO 2002070495 A1 20020912 WO 2002-US6419 20020304

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BE, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KE, LC, LK, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, T2, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG

CA 2440030 A1 20020912 CA 2002-2440030 20020304

US 2002183525 A1 20021205 US 2002-90710 20020304

US 6677458 B2 20040113 US 2003-6677458 20030912

EP 1373229 A1 20040102 EP 2002-717527 20020304

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

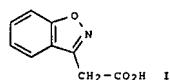
US 2004049053 A1 20040311 US 2003-661109 20030912

PRIORITY APPLN. INFO.: US 2001-273172P P 20010302

US 2001-294847P P 20010531

US 2002-90710 A3 20020304

WO 2002-US6419 W 20020304

OTHER SOURCE(S): CASREACT 137:216942  
GI

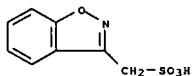
AB A process for the preparation of 1,2-benzisoxazole-3-acetic acid (I) from

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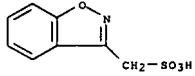
L5 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)  
4-hydroxycoumarin and hydroxylamine.HCl in the presence of a base is disclosed. Compd. I has com. importance as a key intermediate in the prepn. of Zonisamide. For example, a soln. of 4-hydroxycoumarin (100 g), hydroxylamine hydrochloride (150 g) and diethylamine (160 g) in MeOH (500 mL) was heated at reflux for 1 h. The reaction mixt. was evapd. to dryness and the solid dissolved in aq. NaHCO<sub>3</sub> and extd. with ether.

After acidification of the aq. phase, the product was isolated by filtration, washed with water and dried to provide I (99.82 g) in 93 % wt./wt. yield. Advantages of the present invention are: (1) the prep. of I without the use of metallic sodium; and (2) the minimization of reaction side-products, e.g., oxime. The process is thus substantially less hazardous than previous methods. The invention also claims the prep. I or salts of which are converted to 1,2-benzisoxazole-3-methanesulfonamide, i.e., zonisamide.  
IT 73101-64-1P, 1,2-Benzisoxazole-3-methanesulfonic acid sodium salt 342623-49-8P, 1,2-Benzisoxazole-3-methanesulfonic acid 457635-27-7P 457635-28-8P  
RL: IMP (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)  
(product: process for preparation of 1,2-benzisoxazole-3-acetic acid, an intermediate in synthesis of zonisamide)  
RN 73101-64-1 CAPLUS  
CN 1,2-Benzisoxazole-3-methanesulfonic acid, sodium salt (9CI) (CA INDEX NAME)



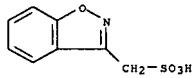
● Na

RN 342623-49-8 CAPLUS  
CN 1,2-Benzisoxazole-3-methanesulfonic acid (9CI) (CA INDEX NAME)



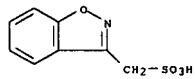
RN 457635-27-7 CAPLUS  
CN 1,2-Benzisoxazole-3-methanesulfonic acid, calcium salt (9CI) (CA INDEX NAME)

L5 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)



● 1/2 Ca

RN 457635-28-8 CAPLUS  
CN 1,2-Benzisoxazole-3-methanesulfonic acid, barium salt (9CI) (CA INDEX NAME)



● 1/2 Ba

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT

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LOGOFF? (Y)/N/HOLD:Y

COST IN U.S. DOLLARS

	SINCE FILE	TOTAL
	ENTRY	SESSION

FULL ESTIMATED COST

13.02	185.33
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

	SINCE FILE	TOTAL
	ENTRY	SESSION

CA SUBSCRIBER PRICE

-1.56	-1.56
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